1 2 Claims: The use of a composition of PKB, its analogues, 3 isoforms, inhibitors, activators and/or the functional 4 equivalents thereof, to regulate glycogen metabolism 5 and/or protein synthesis. 6 7 The use of a composition of PKB, its analogues, 8 isoforms, inhibitors, activators and/or the functional 9 equivalents thereof, for the manufacture of a 10 medicament to regulate glycogen metabolism and/or 11 protein synthesis. 12 13 The use as claimed in claim 1 or claim 2, to 14 combat disease states where glycogen metabolism and/or 15 protein synthesis exhibits abnormality. 16 17 The use as claimed in claim 1, 2 or 3, to combat 18 diabetes. 19 20 The use as claimed in any preceding claim, to 21 22 combat cancer. 23 The use as claimed in claim 5, wherein the cancer 24 is breast, pancreatic or ovarian cancer. 25 26 The use as claimed in any preceding claim, wherein 27 the PKB is PKB α , β or γ , an analogue, isoform, 28 inhibitor, activator or a functional equivalent 29 30 thereof. 31 The use as claimed in any preceding claim, wherein 32 the PKB, its analogue, isoform, or functional 33 equivalent is modified at one or both of amino acids

308 and 473 by phosphorylation and/or mutation.

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- 9 A composition of PKB, its analogues, isoforms,
- 2 inhibitors, activators and/or the functional
- 3 equivalents thereof.

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- 5 10 A peptide having or including the amino acid
- 6 sequence Arg-Xaa-Arg-Yaa-Zaa-Ser/Thr-Hyd, where Xaa is
- 7 any amino acid, Yaa and Zaa are any amino acid, and Hyd
- is a large hydrophobic residue, or a functional
- 9 equivalent of such a peptide.

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- 12 Phe or Leu, or a functional equivalent thereof.

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- 14 12 A peptide as claimed in claim 10 or claim 11,
- wherein Yaa or Zaa or both are an amino acid other than
- 16 glycine.

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- 18 13 A peptide as claimed in claim 10, having the amino
- 19 acid sequence GRPRTSSFAEG, or a functional equivalent
- 20 thereof.

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- 22 14 A method of identifying agents able to influence
- the activity of GSK3, said method comprising:

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- 25 a. exposing a test substance to a substrate of GSK3;
- 26 and
- 27 b. detecting whether said substrate has been
- 28 phosphorylated.

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- 30 15 A method of identifying agents which influence the
- 31 activity of PKB, comprising:

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- 33 a. exposing a test substance to a sample containing
- 34 PKB, to form a mixture;
- 35 b. exposing said mixture to a peptide as claimed in
- 36 claim 10, 11, 12 or 13; and

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| 1 | c. | detecting whether (and, optionally, to what |
|---|----|--|
| 2 | | extent) said peptide has been phosphorylated |

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4 16 A method as claimed in claim 14 or 15, wherein the extent of phosphorylation of the peptide is determined. 5

7 A method as claimed in claim 15, wherein the phosphorylation state(s) of one or both of amino acids 308 and 473 on PKB is determined.

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11 18 A method as claimed in any one of claims 14 to 17, 12 wherein the test substance is an analogue, isoform,

13 inhibitor, or activator of PKB.

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15 19 A method as claimed in any one of claims 14 to 18, 16 wherein steps a or b (or both) are carried out in the 17 presence of divalent cations and ATP.

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A method of treatment of the human or non-human 20 animal body, said method comprising administering PKB, its analogues, inhibitors, stimulators or functional equivalents thereof to said body.

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A method as claimed in claim 20, to combat disease 21 states where glycogen metabolism and/or protein synthesis exhibits abnormality.

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28 22 A method as claimed in claim 20 or 21, to combat diabetes. 29

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31 23 A method as claimed in claim 20 or 22, to combat cancer. 32

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34 24 A method as claimed in claim 23, wherein the 35 cancer is breast, pancreatic or ovarian cancer.

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1 25 A method as claimed in any one of claims 20 to 24, wherein the PKB is PKB α , β or γ , an analogue, isoform, inhibitor, activator or a functional equivalent thereof.

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An agent capable of influencing the activity of PKB, its isoforms, analogues and/or functional equivalents, by modifying amino acids 308 and/or 473 by phosphorylation or mutation.

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11 27 A method of determining the ability of a substance 12 to affect the activity or activation of PKB, the method 13 comprising exposing the substance to PKB and 14 phosphatidyl inositol polyphosphate and determining the 15 interaction between PKB and the phosphatidyl inositol 16 polyphosphate.

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A method of determining the ability of a substance to combat diabetes, cancer, or any disorder which involves irregularity of protein synthesis or glycogen metabolism, the method comprising exposing the substance to PKB and phosphatidyl inositol polyphosphate and determining the interaction between PKB and the phosphatidyl inositol polyphosphate.

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29 A method as claimed in claim 27 or claim 28, wherein the interaction between PKB and the phosphatidyl inositol polyphosphate is measured by assessing the phosphorylation state of PKB.

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31 30 A method as claimed in claim 29, wherein the 32 phosphorylation state of PKB at T308 and/or S473 is 33 assessed.

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35 31 A method of identifying activators or inhibitors 36 of GSK3 comprising exposing the substance to be tested 69

to GSK3 and determining the state of activation of GSK3.

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32 A method as claimed in claim 31 wherein the state of activation of GSK3 is determined by assessing its phosphorylation.

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33 A method of determining the suitability of a test substance for use in combatting diabetes, cancer, or any disorder which involves irregularity of protein synthesis or glycogen metabolism, the method comprising exposing the substance to be tested to GSK3 and determining the state of activation of GSK3.

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- A method for screening for inhibitors or activators of enzymes that catalyse the phosphorylation of PKB, the method comprising exposing the substance to be tested to
 - one or more enzymes upstream of PKB;
 - PKB; and (optionally)
 - nucleoside triphosphate

and determining whether (and optionally to what extent) the PKB has been phosphorylated on T308 and/or S473.